



# KANSAS DRUG UTILIZATION REVIEW NEWSLETTER

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**Fall 2011**

Welcome to the fall 2011 edition of the "Kansas Drug Utilization Review Newsletter," published by Health Information Designs, Inc. (HID). This newsletter is part of a continuing effort to keep the Medicaid provider community informed of important changes in the Kansas Medical Assistance Programs (KMAP).

<u>Helpful Web sites</u>	<u>Helpful Numbers</u>	<u>In This Issue</u>
<b>KMAP Web site</b> <a href="https://www.kmap-state-ks.us/">https://www.kmap-state-ks.us/</a>	<b>KMAP PA Help Desk</b> 1-800-285-4978 <b>ACS PA Help Desk</b> 1-877-475-7567 <b>ACS PA Fax</b> 1-866-246-8512	<b>Provider Customer Service</b> 1-800-933-6593 <b>Beneficiary Customer Service</b> 1-800-766-9012 <b>Pharmacy Help Desk</b> 1-866-405-5200
<b>KDHE-DHCF Web site</b> <a href="http://www.kdheks.gov/hcf/">http://www.kdheks.gov/hcf/</a>		Pradaxa Special Storage and Handling Automated Prior Authorization PDL Update

## Pradaxa Special Storage and Handling

In October 2010, Pradaxa was approved by the U.S. Food and Drug Administration (FDA) with special storage and handling requirements. Pradaxa capsules will hydrolyze over time when exposed to humidity, reducing the effectiveness of the medication due to breakdown of the active ingredient.

Recently, the FDA issued an alert about the importance of proper storage and handling of Pradaxa capsules. The Pradaxa label and Medication Guide explain the storage and handling requirements, but the FDA is concerned that these requirements are not commonly known and are not being followed by patients and pharmacies.

Many patients use medication boxes or organizers to aid them in taking their medications on the appropriate schedule. However, because of the potential for product breakdown and loss of potency, consumers should not store Pradaxa in any container other than the original manufacturer packaging. Additionally, pharmacies should only dispense Pradaxa in the original manufacturer packaging. Pharmacies should never repackage Pradaxa capsules in standard amber vials; keeping the capsules in the manufacturer packaging significantly reduces the product breakdown from moisture.

The manufacturer packaging for Pradaxa contains a 30-day supply with a desiccant in the cap to prevent moisture from breaking down the medication. The current package insert states that the product should be discarded 30 days after the original bottle is opened. However, information currently being reviewed by the FDA indicates that the product maintains potency up to 60 days after the bottle is opened, as long as it is stored in the original bottle and the handling requirements are met. These include closing the cap tightly after each use and keeping the bottle away from excessive moisture, heat, and cold.

### Patient Counseling Information for Storage and Handling:

- Keep Pradaxa capsules in the original bottle to protect the product from moisture.
- Open only one bottle of Pradaxa at a time. Once a bottle is opened, it must be used within 60 days.
- Remove only one capsule from the bottle at the time of use and immediately close the bottle tightly.
- Date the bottle to expire 60 days after opening.
- Do not store Pradaxa capsules in medication boxes or organizers.

### References:

Food and Drug Administration. Pradaxa (dabigatran etexilate mesylate) capsules: Special Storage and Handling Requirements. March 30, 2011. Retrieved from <http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm249005.htm>

PL Detail-Document, Extended Storage Time for Pradaxa (Dabigatran). Pharmacist's Letter/Prescriber's Letter. May 2011.

## Automated Prior Authorization

In August 2011, the Kansas Medical Assistance Program (KMAP) began processing pharmacy point of sale (POS) claims through a new automated prior authorization (PA) system. The new system—SmartPA<sup>SM</sup>—is administered by Affiliated Computer Services, Inc. (ACS). KMAP's pharmacy PAs are processed electronically using preset criteria rules compiled through the collaborative efforts of KMAP stakeholders, including the DUR Board.

When a beneficiary presents a prescription to a pharmacy, the provider submits the claim for adjudication. This involves reviewing the claim against any applicable PA criteria, as well as the beneficiary's previous pharmacy and medical claim history, and returning the outcome to the pharmacy. Most prescriptions are reviewed at the POS in less than a second with no involvement of call center or administrative staff. Pharmacies receive a PA determination for the claim at the time of billing, which allows payment for the drug or directs the pharmacy to contact ACS if additional information is needed. ACS operates a PA call center to process claims that do not meet the automated PA criteria.

In addition to processing requests at the point of sale, PA requests may be submitted to the PA call center by telephone and fax. These requests are finalized within 24 hours of receipt and the provider is notified of the outcome after the review is complete. The provider and beneficiary are also notified in writing if the request is denied.

These changes are designed to streamline the PA process and reduce the amount of time or level of effort required by providers to obtain approval of services.

The KMAP PA unit will continue to process PA requests for the following:

- Medications administered in a physician's office
- Hospice medications not covered by the hospice provider
- Medication requests related to presumptive eligibility
- Non-covered AIDS Drug Assistance Program (ADAP) medications
- Medications billed by an out-of-state pharmacy
  - If related to a traumatic brain injury rehabilitation facility
  - If a child must stay out-of-state following a transplant
- Non-covered KAN Be Healthy medications

Providers who need to request a PA in these situations should continue to contact the KMAP PA Department.

### **KMAP Prior Authorization Department**

**Phone:** 1-800-285-4978 or 785-274-5956 (local)

**Fax:** 1-800-913-2229

**Hours of operation:** Monday through Friday, 7:30 a.m. to 5:30 p.m. CST

### **ACS Prior Authorization Call Center**

**Phone:** 1-877-475-7567

**Fax:** 1-866-246-8512

**Hours of operation:** Monday through Friday, 7:30 a.m. to 6:30 p.m. CST

### **References:**

Xerox Corporation and Affiliated Computer Services, Inc. (2011). Automated Prior Authorization. *Process Claims for Less*, page 1.

Kansas Medical Assistance Program. (2011, July 15). Pharmacy and Professional, Pharmacy Prior Authorization.

Retrieved from <https://www.kmap-state-ks.us/Documents/Content/Bulletins/11080%20-%20Pharmacy%20&%20Pro%20-%20Enhanced%20PA.pdf>.

## Want to Provide Feedback?

The Division of Health Care Finance would like to encourage feedback regarding the new automated prior authorization program. Any feedback can be e-mailed to:

**[PriorAuthorization@kdheks.gov](mailto:PriorAuthorization@kdheks.gov)**

# Preferred Drug List

Below is a list of current preferred agents. A complete list of both preferred and non-preferred agents may be found on the KDHE-DHCF Web site. The Preferred Drug List may be updated at any time; please visit the KDHE-DHCF Web site for the most recent version.

[http://www.kdheks.gov/hcf/pharmacy/pharmacy\\_druglist.html](http://www.kdheks.gov/hcf/pharmacy/pharmacy_druglist.html)

<b>Allergy Agents</b> <b>Non-Sedating Antihistamines</b>	<b>Anti-Infectives</b> <b>Anti-Herpes Virus Agents</b>	<b>Cardiovascular Agents</b> <b>CCBs (Non-Dihydropyridines)</b>	<b>Gout Agents</b> <b>Xanthine Oxidase Inhibitors</b>
Claritin® (loratadine) Zyrtec® (cetirizine)	Valtrex® (valacyclovir) Zovirax® (acyclovir) -Oral Dosage Forms Only	Calan® (verapamil IR) Calan SR® (verapamil SR) Cardizem® (diltiazem IR) Covera HS® (verapamil ER) -Brand Name Only Diltia XT® (diltiazem SR) -& AB Rated Generics Isoptin SR® (verapamil SR) Tiazac® (diltiazem) -& AB Rated Generics Verelan® (verapamil SR)	Zylloprim® (allopurinol)
<b>Analgesics</b> <b>Long-Acting Opioids</b>	<b>Biologic Agents</b> <b>Crohn's Disease</b>	<b>ARB/CCB Combos</b>	<b>Injectables</b> <b>Erythropoiesis Stimulating Agents</b>
Morphine Sulfate ER -Generics Only OxyContin® (oxycodone SR)	*Clinical PA may be required Humira® (adalimumab) Remicade® (infliximab)	<b>Central Nervous System</b> <b>Adjunct Antiepileptics</b>	Procrit® (epoetin alfa) <b>Growth Hormones</b> *Clinical PA is required for all agents
<b>Muscle Relaxants (Skeletal)</b> Flexeril 10mg® (cyclobenzaprine) Parafon Forte DSC® (chlorzoxazone) Robaxin® (methocarbamol) Robaxin-750® (methocarbamol) Robaxinal® (methocarbamol/aspirin)	<b>Adult Rheumatoid Arthritis</b> *Clinical PA may be required Enbrel® (etanercept) Humira® (adalimumab)	Keppra® (levetiracetam) Lyrica® (pregabalin) Neurontin® (gabapentin) Zonagra® (zonisamide)	Genotropin® (somatropin) Genotropin MiniQuick® (somatropin) Saizen® (somatropin) Tev-Tropin® (somatropin)
<b>Muscle Relaxants (Spasticity)</b> Lioresal® (baclofen) Zanaflex® (tizanidine) -Tablets Only	<b>Ankylosing Spondylitis</b> *Clinical PA may be required Enbrel® (etanercept) Humira® (adalimumab)	<b>Non-Benzo Sedative Hypnotics</b>	<b>Nasal Agents</b> <b>Intranasal Antihistamines</b>
<b>Oral NSAIDs</b>	<b>Juvenile Idiopathic Arthritis</b> *Clinical PA may be required Enbrel® (etanercept) Humira® (adalimumab)	<b>Novel Sleep Agents</b>	<b>Intranasal Corticosteroids</b>
Advil® (ibuprofen) Aleve® (naproxen) Anaprox® (naproxen sodium) Anaprox DS® (naproxen sodium) Ansaïd® (flurbiprofen) Arthrotec® (diclofenac/misoprostol) Cataflam® (diclofenac potassium) Clinoril® (sulindac) Daypro® (oxaprozin) EC-Naprosyn® (naproxen) Feldene® (piroxicam) -Brand Name Only Indocin® (indomethacin) Lodine® (etodolac) Meclomen® (meclofenamate) Mobic® (meloxicam) Motrin® (ibuprofen) Motrin IB® (ibuprofen) Nalfon® (fenoprofen) Naprelan® (naproxen sodium) Naprosyn® (naproxen) Orudis® (ketoprofen) Orudis KT® (ketoprofen) Oruvail® (ketoprofen) Ponstel® (mefenamic acid) Toradol® (ketorolac) -Limit 5 Day Supply Tolectin DS® (tolmetin) Tolectin 600® (tolmetin) Voltaren® (diclofenac sodium) Voltaren XR® (diclofenac sodium)	<b>Plaque Psoriasis</b> *Clinical PA may be required Humira® (adalimumab) Remicade® (infliximab)	<b>Diabetic Agents</b> <b>Alphaglucoosidase Inhibitors</b>	Flonase® (fluticasone) Nasonex® (mometasone) Veramyst® (fluticasone)
<b>Topical NSAIDs</b> Voltaren® (diclofenac sodium, topical)	<b>Psoriatic Arthritis</b> *Clinical PA may be required Humira® (adalimumab) Remicade® (infliximab)	<b>Biguanides</b>	<b>Ophthalmic Agents</b> <b>Ophthalmic Antihistamine/Mast Cell Stabilizer Combos</b>
<b>Triptans</b> Amerge® (naratriptan) Imitrex® (sumatriptan) -including all generic dosage forms Maxalt® (rizatriptan) Maxalt MLT® (rizatriptan) Relpax® (eletriptan)	<b>Ulcerative Colitis</b> *Clinical PA may be required Remicade® (infliximab)	<b>Insulin (Delivery Systems)</b>	Alaway® (ketotifen) Refresh® (ketotifen) Zaditor® (ketotifen)
<b>Antihyperlipidemics</b> <b>Fibric Acid Derivatives</b> Fenofibrate -Generics Only Lipid® (gemfibrozil) TriCor® (fenofibrate)	<b>Cardiovascular Agents</b> <b>ACE Inhibitors</b>	<b>Long-Acting Insulins</b>	<b>Ophthalmic Prostaglandin Analogs</b>
<b>Statins</b> Lipitor® (atorvastatin) Zocor® (Simvastatin)	<b>ARBs</b>	<b>DPP-4 Inhibitors</b>	<b>Ophthalmic NSAIDs</b>
	Cozaar® (losartan/HCTZ) Diovan® (valsartan) Diovan HCT® (valsartan/HCTZ) Hyzaar® (losartan) Micardis® (telmisartan) Micardis HCT® (telmisartan/HCTZ)	<b>Meglitinides</b>	<b>Osteoporosis Agents</b> <b>Bisphosphonates</b>
	<b>Beta-Blockers</b>	<b>Insulin (Delivery Systems)</b>	Fosamax® (alendronate) Fosamax Plus D® (alendronate/cholecalciferol)
	Betapace® (sotalol) Betapace AF® (sotalol AF) Blocadren® (timolol) Corgard® (nadolol) Coreg® (carvedilol) Coreg CR® (carvedilol CR) Inderal® (propranolol) InnoPran XL® (propranolol XL) Kerlone® (betaxolol) Lopressor® (metoprolol tartrate) Propranolol Intensol® (propranolol) Sectral® (acebutolol) Tenormin® (atenolol) Toprol XL® (metoprolol succinate) Viskin® (pindolol)	<b>Long-Acting Insulins</b>	<b>Respiratory</b> <b>Inhaled Corticosteroids</b>
	<b>CCBs (Dihydropyridines)</b>	<b>2<sup>nd</sup> Generation Sulfonylureas</b>	Flovent Diskus® (fluticasone) Flovent HFA® (fluticasone) Pulmicort Respules® (budesonide) -6 & Under Only QVAR® (beclomethasone)
	Adalat CC® (nifedipine ER) Cardene® (nicardipine IR) DynaCirc® (isradipine IR) DynaCirc CR® (isradipine CR) Norvasc® (amlodipine) Procardia XL® (nifedipine ER)	<b>Thiazolidinediones</b>	<b>Inhaled Long Acting Inhaled Beta<sub>2</sub> Agonists</b>
	<b>ACE Inhibitor/CCB Combos</b> Lotrel® (benazepril/amlodipine)	<b>Gastrointestinal Agents</b> <b>Serotonin 5HT<sub>3</sub> Antagonists</b>	<b>Inhaled Long Acting Inhaled Beta<sub>2</sub> Agonists/Corticosteroid Combos</b>
		<b>Pancreatic Enzyme Replacements</b>	Advair® (fluticasone/salmeterol) Advair HFA® (fluticasone/salmeterol) Dulera® (formoterol/mometasone) Symbicort® (budesonide/formoterol)
		<b>Proton Pump Inhibitors</b>	<b>Inhaled Short Acting Inhaled Beta<sub>2</sub> Agonists</b>
		<b>H<sub>2</sub> Antagonists</b>	ProAir HFA® (albuterol) Proventi® (albuterol) Ventolin® (albuterol) Ventolin HFA® (albuterol)
		<b>Anticholinergics</b>	<b>Urologic Agents</b> <b>Anticholinergics</b>
		Dexilant® (dexlansoprazole) Prevacid® (lansoprazole) Prevacid SoluTab® (lansoprazole) Prilosec® (omeprazole)	Detrol® (tolterodine) Detrol LA® (tolterodine LA) Ditropan® (oxybutynin) Toviaz® (fesoterodine) Vesicare® (solifenacin)



Health Information Designs, Inc. (HID) was founded in 1976 with a mission to provide drug utilization review (DUR) services for state Medicaid agencies. In 1997, HID was acquired by HDI Solutions and subsequently has experienced strong and steady growth

as a premium healthcare and pharmacy support services provider.

Currently, HID works with government agencies in 27 states, including Medicaid agencies and Boards of Pharmacy. HID's efforts in these states monitor, manage or administer more than one-third of the nation's Medicaid budget. The work performed by HID has a daily impact on the healthcare of more than 66 million Americans.

HID currently lists among our clients 17 state Medicaid programs, 14 state health department programs, and several commercial pharmacy benefit management (PBM) organizations. To serve this geographically-widespread client base, in addition to our home offices in Auburn, Alabama, we have staff in Arkansas, Connecticut, Kansas, Maryland, Mississippi, Texas and Wisconsin.

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